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INTERNATIONAL PRELIMINARY REPORT ON PATENT
(Chapter II of the Patent Cooperation Treaty)

DUE DATE:	—
FORMALITIES:	CA RP
PAT. OFF:	AK
ON DB FABRI TV	15/6/06
CASE NO:	PV0407-R

(PCT Article 36 and Rule 70)

Applicant's or agent's file reference PU0407-PCT ~	FOR FURTHER ACTION See Form PCT/IPEA/416	
International application No. PCT/SE2005/000229 ~	International filing date (day/month/year) 21-02-2005 ~	Priority date (day/month/year) 26-02-2004 ~
International Patent Classification (IPC) or national classification and IPC See Supplemental Box		
Applicant GE HEALTHCARE BIO-SCIENCES AB et al		

- This report is the international preliminary examination report, established by this International Preliminary Examining Authority under Article 35 and transmitted to the applicant according to Article 36.
- This REPORT consists of a total of 6 sheets, including this cover sheet.
- This report is also accompanied by ANNEXES, comprising:
 - ☐ (sent to the applicant and to the International Bureau) a total of _____ sheets, as follows:
 - ☐ sheets of the description, claims and/or drawings which have been amended and are the basis of this report and/or sheets containing rectifications authorized by this Authority (see Rule 70.16 and Section 607 of the Administrative Instructions).
 - ☐ sheets which supersede earlier sheets, but which this Authority considers contain an amendment that goes beyond the disclosure in the international application as filed, as indicated in item 4 of Box No. I and the Supplemental Box.
 - ☐ (sent to the International Bureau only) a total of (indicate type and number of electronic carrier(s)) _____, containing a sequence listing and/or tables related thereto, in electronic form only, as indicated in the Supplemental Box Relating to Sequence Listing (see Section 802 of the Administrative Instructions).

- This report contains indications relating to the following items:

- | | | |
|-------------------------------------|--------------|---|
| <input checked="" type="checkbox"/> | Box No. I | Basis of the report |
| <input type="checkbox"/> | Box No. II | Priority |
| <input type="checkbox"/> | Box No. III | Non-establishment of opinion with regard to novelty, inventive step and industrial applicability |
| <input type="checkbox"/> | Box No. IV | Lack of unity of invention |
| <input checked="" type="checkbox"/> | Box No. V | Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability, citations and explanations supporting such statement |
| <input type="checkbox"/> | Box No. VI | Certain documents cited |
| <input type="checkbox"/> | Box No. VII | Certain defects in the international application |
| <input checked="" type="checkbox"/> | Box No. VIII | Certain observations on the international application |

Date of submission of the demand 01-09-2005	Date of completion of this report 29-05-2006
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Form PCT/IPEA/409 (cover sheet) (April 2005)

INTERNATIONAL PRELIMINARY REPORT ON PATENTABILITY

International application No.

PCT/SE2005/000229

Supplemental Box

In case the space in any of the preceding boxes is not sufficient.

Continuation of: **Cover sheet**

International patent classification (IPC)

C12N15/10 (2006.01)

B01D 15/08 (2006.01)

INTERNATIONAL PRELIMINARY REPORT ON PATENTABILITY

International application No.

PCT/SE2005/000229

Box No. I Basis of the report

1. With regard to the language, this report is based on:

- ☒ the international application in the language in which it was filed
- ☐ a translation of the international application into _____, which is the language of a translation furnished for the purposes of:
- ☐ international search (Rules 12.3(a) and 23.1(b))
 - ☐ publication of the international application (Rule 12.4(a))
 - ☐ international preliminary examination (Rules 55.2(a) and/or 55.3(a))

2. With regard to the elements of the international application, this report is based on *(replacement sheets which have been furnished to the receiving Office in response to an invitation under Article 14 are referred to in this report as "originally filed" and are not annexed to this report)*:

- ☒ the international application as originally filed/furnished
- ☐ the description:
- pages _____ as originally filed/furnished
- pages* _____ received by this Authority on _____
- pages* _____ received by this Authority on _____
- ☐ the claims:
- pages _____ as originally filed/furnished
- pages* _____ as amended (together with any statement) under Article 19
- pages* _____ received by this Authority on _____
- pages* _____ received by this Authority on _____
- ☐ the drawings:
- pages _____ as originally filed/furnished
- pages* _____ received by this Authority on _____
- pages* _____ received by this Authority on _____
- ☐ a sequence listing and/or any related table(s) – see Supplemental Box Relating to Sequence Listing.

3. ☐ The amendments have resulted in the cancellation of:

- ☐ the description, pages _____
- ☐ the claims, Nos. _____
- ☐ the drawings, sheets/figs _____
- ☐ the sequence listing (*specify*): _____
- ☐ any table(s) related to the sequence listing (*specify*): _____

4. ☐ This report has been established as if (some of) the amendments annexed to this report and listed below had not been made, since they have been considered to go beyond the disclosure as filed, as indicated in the Supplemental Box (Rule 70.2(c)).

- ☐ the description, pages _____
- ☐ the claims, Nos. _____
- ☐ the drawings, sheets/figs _____
- ☐ the sequence listing (*specify*): _____
- ☐ any table(s) related to the sequence listing (*specify*): _____

* If item 4 applies, some or all of those sheets may be marked "superseded."

INTERNATIONAL PRELIMINARY REPORT ON PATENTABILITY

International application No.

PCT/SE2005/000229

Box No. V Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

1. Statement

Novelty (N)	Claims	<u>7-8.16</u>	YES
	Claims	<u>1.2.11.12</u>	NO
Inventive step (IS)	Claims		YES
	Claims	<u>1-6.9-15.17-22</u>	NO
Industrial applicability (IA)	Claims	<u>1-22</u>	YES
	Claims		NO

2. Citations and explanations (Rule 70.7)

The invention relates to methods for the isolation of plasmids using a separation matrix with anion exchange groups. The chosen pore size distribution does not allow access of plasmids to the pore surfaces.

The most relevant documents cited in the International Search Report are:

D1: WO9963076A1
D2: WO0137987A1
D3: US6270970B1

Document D1 discloses a method of purifying plasmids using a TMAE anion exchange chromatographic column (see claims 1-3). The used matrix is a fractogel TMAE anion exchange resin. These resins are known to have particle sizes between 20-40 μm for TMAE S and 40-90 μm for TMAE M. The pore size is about 800 Å (see Merck website).

Thus, D1 is considered to disclose a method of isolating plasmids with the steps of providing a separation matrix comprised of porous carriers, which carrier present anion exchange groups on external surfaces as well as pore surfaces and a pore size distribution that does not allow access of plasmids to pore surfaces; contacting said matrix with a liquid to absorb plasmids to ligands present on the external surfaces of the separation matrix

.../...

Supplemental Box

In case the space in any of the preceding boxes is not sufficient.

Continuation of: V

Document D2 discloses separation methods for plasmids. In example 3 a separation of plasmids is performed with anion exchange chromatography. The plasmids are bound to the separation medium B and its charged outer surfaces of the anion-exchanger. It is not stated that the plasmids have access to the pores.

Document D3 relates to mixed-bed solid phases for isolation of nucleic acids such as plasmids. The solid phase of the different beds comprise magnetic silica particles (particle size below 15 μm), see column 12. The solid phase can be with or without pores with size sufficiently large to admit the target nucleic acid in to the interior of the particles. The anion exchanger phase can be Sepharose but is not limited thereto.

With background of D1-D3, and as a consequence of unclear claims (see box VIII), the method according to claim 1 and the use according to claim 11 lacks novelty. Further, the DNA exclusion limits covered by D1-D3 are assumed to be at least about 270 base pairs. Therefore, also claims 2 and 12 lacks novelty.

The claims 3-6, 9-10, 13-15 and 17 are considered to involve particular detail executions obvious to a person skilled in the art. Therefore, the invention according to these claims is not considered to involve an inventive step.

It is also considered to be obvious to a person skilled in the art to develop a kit for the method described in D1 or D2. Therefore the invention according to claims 18-22 lacks an inventive step.

Claims 7-8 and 16 are novel and considered to involve an inventive step.

Box No. VIII Certain observations on the international application

The following observations on the clarity of the claims, description, and drawings or on the question whether the claims are fully supported by the description, are made:

Claims 1, 11 and 18 do not meet the requirements of Article 6 PCT in that the matter for which protection is sought is not clearly defined. The claims attempts to define the subject-matter in terms of the result to be achieved (...pore size distribution that does not allow access of plasmids...) which merely amounts to a statement of the underlying problem. The technical features necessary for achieving this result should be added.

In claims 2, 12 and 19 the matrix is characterised by a DNA exclusion limit of at least about 270 base pairs. This way of characterising a matrix is known in the field but is not a common way of defining and comparing gels. Further, the limit of "about" 270 base pairs is unclear (see PCT GL 5.38).

Claims 1-2, 11-12 and 18-19 have been drafted as separate independent claims of the same category. They appear to relate effectively to the same subject-matter and to differ from each other only with regard to the choice of specific words. The aforementioned claims therefore lack conciseness. See PCT Article 6 and 5.42 Guidelines.